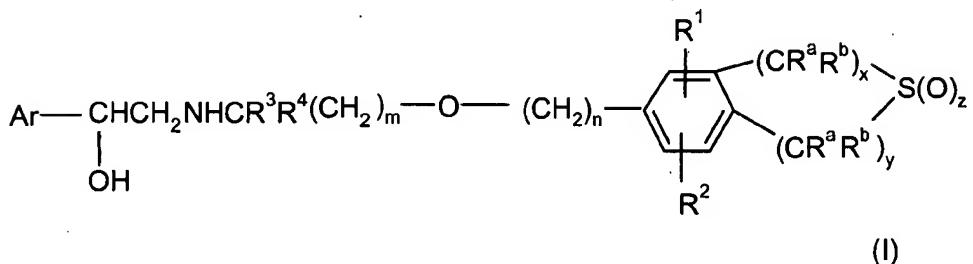


Amendments To The Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

What is claimed is:

1. (Currently Amended) A compound of formula (I)



or a salt, solvate, or physiologically functional derivative thereof, wherein:

m is an integer of from 2 to 8;

n is an integer of from 3 to 11;

with the proviso that m + n is 5 to 19;

x is zero and y is an integer of 2 or 3 or

y is zero and x is an integer of 2 or 3;

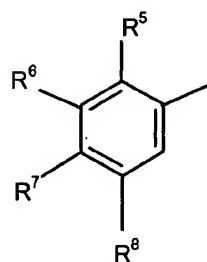
z is zero or an integer of 1 or 2;

R^a and R^b are independently selected from hydrogen and C₁₋₄alkyl;

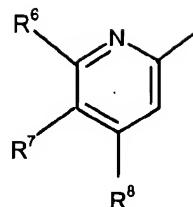
R¹ and R² are independently selected from hydrogen, C₁₋₆alkyl, C₁₋₆alkoxy, halo, phenyl, and C₁₋₆haloalkyl;

R³ and R⁴ are independently selected from hydrogen and C₁₋₄alkyl with the proviso that the total number of carbon atoms in R³ and R⁴ is not more than 4;

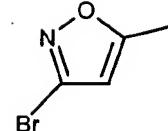
Ar is a group selected from



(a)

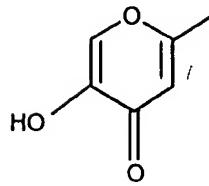


(b)



(c)

and



(d)

wherein R⁸ represents hydrogen, halogen, -(CH₂)_qOR⁹, -NR⁹C(O)R¹⁰, -NR⁹SO₂R¹⁰, -SO₂NR⁹R¹⁰, -NR⁹R¹⁰, -OC(O)R¹¹ or -OC(O)NR⁹R¹⁰, and R⁵ represents hydrogen, halogen or C₁₋₄alkyl;

or R⁶ represents -NHR¹² and R⁵ and -NHR¹² together form a 5- or 6-membered heterocyclic ring;

R⁷ represents hydrogen, halogen, -OR⁹ or -NR⁹R¹⁰;

R⁸ represents hydrogen, halogen, haloC₁₋₄ alkyl, -OR⁹, -NR⁹R¹⁰, -OC(O)R¹¹ or -OC(O)NR⁹R¹⁰;

R⁹ and R¹⁰ independently represent hydrogen or C₁₋₄ alkyl or R⁹ and R¹⁰ together with the nitrogen atom to which they are attached form a 5-, 6- or 7-membered nitrogen-containing ring,

R¹¹ represents an aryl (eg phenyl or naphthyl) group which may be unsubstituted or substituted by one or more substituents selected from halogen, C₁₋₄ alkyl, hydroxy, C₁₋₄ alkoxy or halo C₁₋₄ alkyl; and

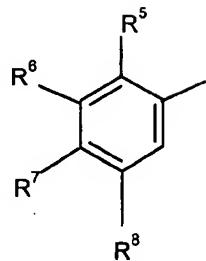
q is zero or an integer from 1 to 4.

2. (Original) A compound according to claim 1 wherein R³ and R⁴ are independently selected from hydrogen and methyl.

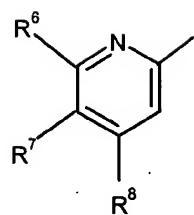
3. (Previously Presented) A compound according to claim 1 wherein R¹ and R² each represent hydrogen.

4. (Previously Presented) A compound according to claim 1 wherein the integer m is 4, 5 or 6 and n is 3, 4, 5 or 6.

5. (Previously Presented) A compound according to claim 1 wherein the group Ar is selected from groups (a) and (b).

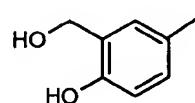


(a)

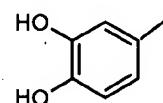


(b)

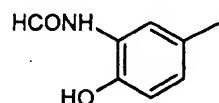
6. (Previously Presented) A compound according to claim 5 wherein groups (a) and (b) are selected from the group consisting of (i) to (xxi):



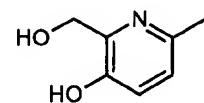
(i)



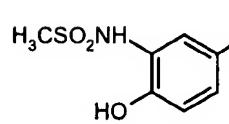
(ii)



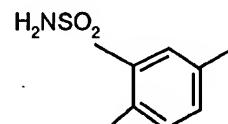
(iii)



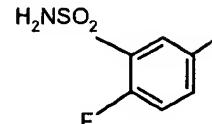
(iv)



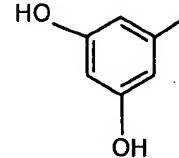
(v)



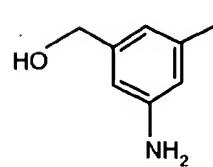
(vi)



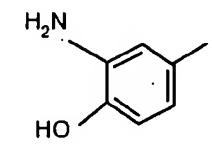
(vii)



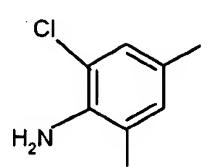
(viii)



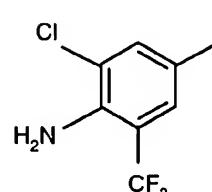
(ix)



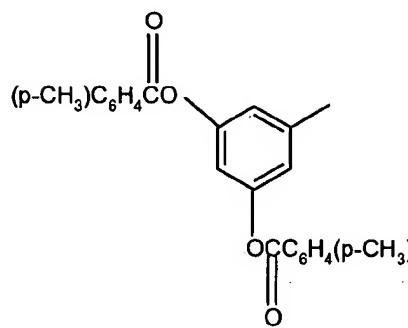
(x)



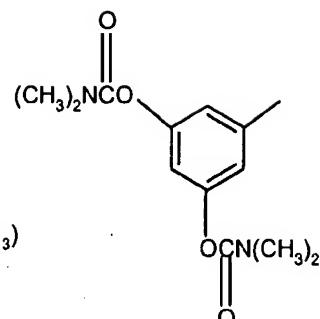
(xi)



(xii)



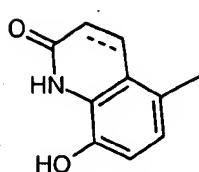
(xiii)



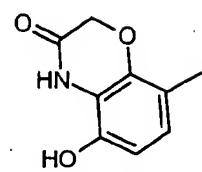
(xiv)



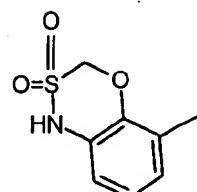
(xv)



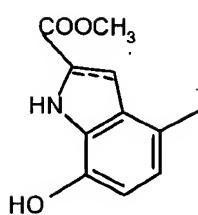
(xvi)



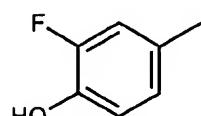
(xvii)



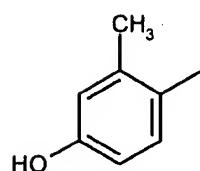
(xviii)



(xix)

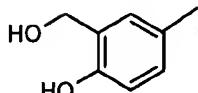


(xx)



(xxi)

7. (Previously Presented) A compound of formula (I) according to claim 6 wherein Ar represents group (i).



(i)

8. (Previously Presented) A compound of formula (I) according to claim 1 wherein z represents 2.

9. (Previously Presented) A compound of formula (I) according to claim 1 which is selected from the group consisting of:

4-[(1*R*)-2-({6-[4-(1,1-Dioxido-2,3-dihydro-1-benzothien-6-yl)butoxy]hexyl}amino)-1-hydroxyethyl]-2-(hydroxymethyl)phenol;
4-[(1*r*)-2-({6-[4-(1,1-dioxido-3,4-dihydro-2*h*-thiochromen-7-yl)butoxy]hexyl}amino)-1-hydroxyethyl]-2-(hydroxymethyl)phenol;

salts thereof, solvates thereof and physiologically functional derivatives thereof.

10. (Previously Presented) A method for the prophylaxis or treatment of a clinical condition in a mammal for which a selective β₂-adrenoreceptor agonist is indicated, which comprises administating a therapeutically effective amount of a compound of formula (I), according to claim 1, or a pharmaceutically acceptable salt, solvate, or physiologically functional derivative thereof.

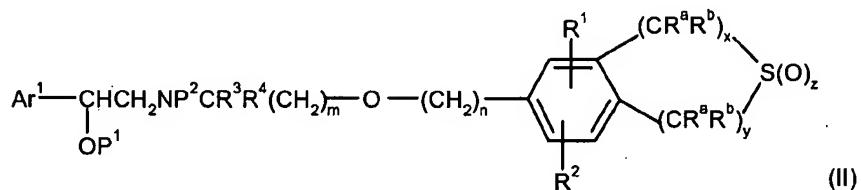
11-12. (Canceled)

13. (Previously Presented) A pharmaceutical formulation comprising a compound of formula (I), according to claim 1, or a pharmaceutically acceptable salt, solvate, or physiologically functional derivative thereof, and a pharmaceutically acceptable carrier or excipient, and optionally one or more other therapeutic ingredients.

14. (Cancelled)

15.

(Previously Presented) A process for the preparation of a compound of formula (I), according to claim 1, or a salt, solvate, or physiologically functional derivative thereof, which comprises:
deprotecting a protected intermediate of formula (II):



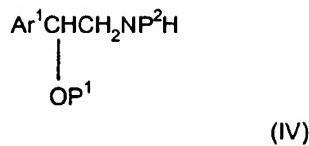
or a salt or solvate thereof, wherein R^a, R^b, R¹, R², R³, R⁴, m, n; x, y and z are as defined for the compound of formula (I) or (Ia), Ar¹ represents an optionally protected form of Ar; and P¹ and P² are each independently either hydrogen or a protecting group, such that the compound of formula (II) contains at least one protecting group

wherein said deprotecting step is optionally followed by one or more of the following steps in any order selected from the group consisting of:

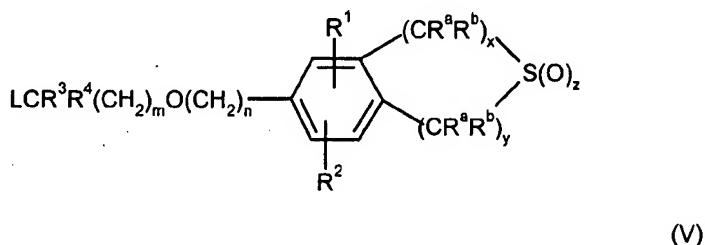
- (i) removing any protecting groups;
- (ii) separating an enantiomer from a mixture of enantiomers;
- (iii) converting one compound of formula (I) to a different compound of formula (I); and
- (iv) converting the product to a corresponding salt, solvate, or physiologically functional derivative thereof.

16.

(Previously Presented) A process for the preparation of a compound of formula (I), according to claim 1, or a salt, solvate, or physiologically functional derivative thereof, which comprises reacting a compound of formula (IV):



wherein Ar^1 represents an optionally protected form of Ar; and P^1 and P^2 each independently represent hydrogen or a protecting group, with a compound of formula (V):



wherein L is a leaving group, and R^a , R^b , R^1 , R^2 , R^3 , R^4 , n, m, x, y and z are as defined for compounds of formula (I);

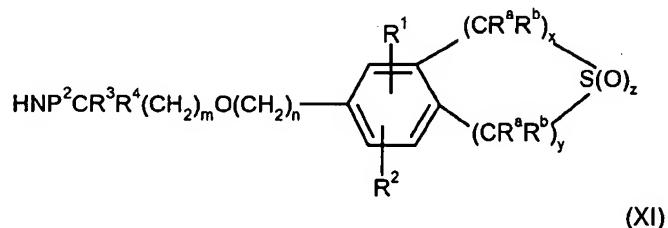
wherein said reacting step is optionally followed by one or more of the following steps in any order selected from the group consisting of:

- (i) removing any protecting groups;
- (ii) separating an enantiomer from a mixture of enantiomers;
- (iii) converting one compound of formula (I) to a different compound of formula (I); and
- (iv) converting the product to a corresponding salt, solvate, or physiologically functional derivative thereof.

17. (Previously Presented) A process for the preparation of a compound of formula (I), according to claim 1, or a salt, solvate, or physiologically functional derivative thereof, which comprises reacting a compound of formula (X):



wherein Ar¹ represents an optionally protected form of Ar; P¹ independently represents hydrogen or a protecting group and L is a leaving group, with an amine of formula (XI):



wherein R^a, R^b, R¹, R², R³, R⁴, m, n, x, y and z are as defined; and P² represents hydrogen or a protecting group;

wherein said reacting step is optionally followed by one or more of the following steps in any order selected from the group consisting of:

- (i) removing any protecting groups;
- (ii) separating an enantiomer from a mixture of enantiomers;
- (iii) converting one compound of formula (I) to a different compound of formula (I); and
- (iv) converting the product to a corresponding salt, solvate, or physiologically functional derivative thereof.

18. (Previously Presented) The method according to claim 10, wherein the mammal is a human.

16
19. (Previously Presented) The process according to Claim **16**, wherein
L is a halo or sulfonate leaving group.

¹⁷
~~20.~~ (Previously Presented) The process according to Claim ¹⁶~~19~~, wherein L is selected from the group consisting of an alkylsulfonate, an aryl sulfonate, and a haloalkylsulfonate.

¹⁸
~~21.~~ (Previously Presented) The process according to Claim ¹⁴~~17~~, wherein L is a halo or sulfonate leaving group.

¹⁹
~~22.~~ (Previously Presented) The process according to Claim ¹⁸~~21~~, wherein L is selected from the group consisting of an alkylsulfonate, an aryl sulfonate, and a haloalkylsulfonate.